

# BGMUT: NCBI dbRBC database of allelic variations of genes encoding antigens of blood group systems

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## ABSTRACT

Analogous to human leukocyte antigens, blood group antigens are surface markers on the erythrocyte cell membrane whose structures differ among individuals and which can be serologically identified. The Blood Group Antigen Gene Mutation Database (BGMUT) is an online repository of allelic variations in genes that determine the antigens of various human blood group systems. The database is manually curated with allelic information collated from scientific literature and from direct submissions from research laboratories. Currently, the database documents sequence variations of a total of 1251 alleles of all 40 gene loci that together are known to affect antigens of 30 human blood group systems. When available, information on the geographic or ethnic prevalence of an allele is also provided. The BGMUT website also has general information on the human blood group systems and the genes responsible for them. BGMUT is a part of the dbRBC resource of the National Center for Biotechnology Information, USA, and is available online at <http://www.ncbi.nlm.nih.gov/projects/gv/rbc/xslcgi.fcgi?cmd=bgmut>. The database should be of use to members of the transfusion medicine community, those interested in studies of genetic variation and related topics such as human migrations, and students as well as members of the general public.

## INTRODUCTION

The BGMUT Blood Group Antigen Gene Mutation Database documents variations in genes that encode antigens for human blood groups. It is a part of the dbRBC resource (1) of the National Center for

Biotechnology Information (NCBI) of USA and can be freely accessed online at <http://www.ncbi.nlm.nih.gov/projects/gv/rbc/xslcgi.fcgi?cmd=bgmut>.

Recent documentation of the extent and the surprisingly high numbers of mutations in the human genome have suggested that, perhaps with the exception of identical twins, no two individuals bear exact copies of chromosomal DNA. In those studies, DNA of random subjects is compared but more often phenotypic differences observed in disease states, whether single gene inherited disorders or in association studies of complex conditions are taken as criteria for selection of individuals whose DNA is examined for sequence changes. In the latter studies, large fragments of DNA are usually examined and compared statistically to matched control individuals. Changes in blood group phenotypes are another criterion for selection of subjects who may show differences in sequences of two or more defined sets of genes. These genes encode a group of red cell membrane proteins that are polymorphic in the population and are defined as blood group antigens; in addition, these genes may encode certain glycosyltransferases that are involved in the synthesis of red cell membrane glycans whose structures also differ among individuals. The former group consists of structural molecules, channels, adhesion molecules or enzymes and, excluding their red cell membrane location, can be considered as representative of any human protein, whereas the latter are similar to most other glycosyltransferases. Evidence suggests that sequence changes in these proteins or even their absence, such as in null phenotypes, is not, in most cases, physiologically harmful.

The proteins or the glycans fulfill their role as blood group antigens because they are polymorphic in the population and their sequence changes can be readily predicted by serological approaches; they are known as ‘antigens’ because in the course of transfusion, or pregnancy, the presence of a variant protein epitope is recognized as ‘non-self’ and may ultimately result in an adverse

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immunological reaction. The use of transfusion being ubiquitous in the practice of medicine, populations worldwide are serologically tested and variant antigens and their genes, in contrast to many other variant genes are being documented in a large number of diverse populations. Although some variants occur rarely and, may only be observed in a single individual or family, others appear in unexpectedly large populations, such as the MiIII phenotype encoded by the MiIII *GYPA* gene, in Taiwan (incidence can be as high as 88% among Ami tribes) (2). For many alleles, the database also provides information on the geographic or ethnic origin of alleles and/or their associated serological phenotypes when such information was presented in the publications describing the alleles. This may be of use to those interested in population migrations.

## HISTORY AND CURRENT STATE

BGMUT was developed in 1999 as a locus-specific gene mutation database under the aegis of the Human Variation Genome Society. It was curated under the direction of one of the authors (OOB) with original information contributed by more than a dozen blood group system experts. The database was hosted online by the Department of Biochemistry at the Albert Einstein College of Medicine, New York. BGMUT was identified as one of three model locus-specific databases from more than 200 in a scholarly review (3). In 2006, BGMUT became a part of the dbRBC resource of the NCBI. At dbRBC, curatorship and direction for maintenance of the database has been provided by another of the authors (WH). The number of alleles in BGMUT has approximately doubled since 2004 when the database was first described in a scholarly publication (4). This publication has been referred to many times in the scientific literature indicating that BGMUT has been a useful resource. Links to BGMUT database records are available on relevant pages of the Wikipedia online encyclopedia, and on many of NCBI's online resources. BGMUT is also a part of the PhenCode project which attempts to integrate genetic variation data with the UCSC Genome Browser (5).

As of August 2011, BGMUT had 1251 alleles belonging to 40 genes that are together responsible for 30 human blood group systems (Table 1). Alleles of some genes, such as *ABO* and *RHCE/D* which are, respectively, responsible for the ABO and Rh blood group systems most frequently examined in the populations, are more numerous than those of others (Table 1). As per the International Society for Blood Transfusion, there are 30 human blood group systems (<http://ibgrl.blood.co.uk/ISBT%20Pages/ISBT%20Terminology%20Pages/Table%20of%20blood%20group%20systems.htm>), all of which are covered by BGMUT. The Globoside (GLOB) system is currently considered a part of the P1PK system in BGMUT, which additionally considers the system of T and Tn antigens as a separate blood group system.

## DATABASE ARCHITECTURE

BGMUT is accessible online for view, search or for administration as a website in the form of HTML (hyper-text markup language) pages (Figures 1 and 2). A Microsoft SQL Server relational database is used for data storage. Programmatic code in SQL (structured query language), C++ and XSLT (extensible structured language transformations) languages is used for interaction with the SQL Server database and for rendering BGMUT's web-based interface. Raw data on all or a subset of the alleles described in BGMUT can be downloaded as tab-delimited or comma-separated (CSV) text formats from the BGMUT website. Compilations of allelic sequences for the ABO, H, MNS and Rh systems in the Microsoft Excel format are also available for download.

## DATABASE CURATION AND ALLELE SUBMISSION

The BGMUT database is curated manually. Allelic information is periodically collated from scientific literature, as is the case for a majority of the alleles listed in BGMUT, or is obtained as direct submissions from researchers through the database website. During the process of curation, good quality of methods used in a study is ascertained. The new candidate allele's sequence that has been published and/or submitted to a publicly available repository such as NCBI's GenBank is compared to BGMUT's reference allele for the gene sequence. The sequence positions and the kinds of the deduced amino acid changes are also verified. Authors are consulted in case of a question or disagreement. For direct submission of information on a new allele for inclusion in the database, a scientific publication describing the allele is not required. However, submitters are encouraged to deposit the allele's sequence in a publicly available repository. In the absence of a scientific publication, this is a requirement.

## ALLELES IN BGMUT

Alleles in BGMUT are grouped by the blood group system that the genes they belong to affect. For each allele in the database, BGMUT provides details on the nucleotide changes and the deduced amino acid changes in the protein encoded by the gene the allele belongs to. These changes are in context of a 'reference' allele that itself is included in BGMUT, and is the same for all alleles of a gene. Besides the information on the sequence changes, BGMUT also details for an allele the frequency of occurrence, the associated blood group phenotype, references to the studies that identified and characterized the allele and accession numbers of the relevant sequences in NCBI GenBank when such information is available. GenBank accession numbers, however, are not available for many alleles because though they have been described in published literature, their sequences were not deposited in the repository by the authors. When known, the regions of the gene or cDNA that were sequenced to identify the allele, the prevalence of the allele in different geographical

**Table 1.** Blood group systems in the BGMUT database<sup>a</sup>

System (symbol)	Symbol	Chromosomal location	Genes	
			Nature of encoded protein	Alleles
ABO (ABO)	ABO	9q34.2	Galactosyltransferase and <i>N</i> -acetylgalactosaminyltransferase	272
Chido/Rodgers (CH/RG)	C4A	6p21.3	Complement factor	3
	C4B	6p21.3	Complement factor	4
Colton (CO)	AQP1	7p14	Water channel	11
Cromer (CROM)	CD55	1q32	Complement cascade regulator	15
Diego (DI)	SLC4A1	17q21-q22	Anion exchanger	91
Dombrock (DO)	ART4	12p13-p12	ADP ribosyltransferase	20
Duffy (FY)	DARC	1q21-q22	Chemokine receptor	11
Gerbich (GE)	GPC	2q14-q21	Cytoketal element	10
Gill (GIL)	AQP3	9p13	Water channel	8
H (H)	FUT1	19q13.3	Fucosyltransferase	45
	FUT2	19q13.3	Fucosyltransferase	56
I (I)	GCNT2	6p24.2	<i>N</i> -acetylglucosaminyltransferase	13
Indian (IN)	CD44	11p13	Adhesion molecule	4
John Milton Hagen (JMH)	SEMA7A	15q22.3-q23	Adhesion and signaling molecule?	12
Kell (KEL)	KEL	7q33	Metalloendopeptidase?	57
Kidd (JK)	SLC14A1	18q11-q12	Urea transporter	18
Knops (KN)	CR1	1q32	Cell surface receptor	32
Kx (XK)	XK	Xp21.1	Membrane transport protein?	35
Landsteiner-Weiner (LW)	ICAM4	19p13.2-cen	Adhesion molecule	4
Lewis (LE) <sup>b</sup>	FUT3	19p13.3	Fucosyltransferase	52
	FUT6	19p13.3	Fucosyltransferase	20
	FUT7	9q34.3	Fucosyltransferase	2
Lutheran (LU)	BCAM	19q13.2	Adhesion molecule	20
MNS (MNS)	GYPA	4q31.21	Unknown	38 <sup>c</sup>
	GYPB	4q31.21	Unknown	35 <sup>c</sup>
	GYPE	4q31.1	Unknown	2 <sup>c</sup>
Ok (OK)	BSG	19p13.3	Plasma membrane protein	5
P1PK (P1PK) including globoside (GLOB)	B3GALNT1	3q25	<i>N</i> -acetylgalactosaminyltransferase	9
	A4GALT	22q13.2	Galactosyltransferase	37
Raph (RAPH)	CD151	11p15.5	Adhesion molecule	4
Rh (RH)	RHCE	1p36.11	Unknown	103
	RHD	1p36.11	Unknown	185
Rh-associated glycoprotein (RHAG)	RHAG	6p21.1-p11	Ammonium and carbon dioxide transporter	23
Scianna (SC)	ERMAP	1p34.2	Adhesion molecule?	9
T/Tn	C1GALT1C1	Xq24	Galactosyltransferase	8
	C1GALT1	7p21.3	Galactosyltransferase	1
Xg (XG)	XG	Xp22.33	Unknown	1
	CD99	Xp22.32, Yp11.3	Adhesion molecule	1
Yt (YT)	ACHE	7q22	Acetylcholinesterase	4

<sup>a</sup>As of August 2011; total 30 systems, 40 genes and 1251 alleles.<sup>b</sup>The FUT3, FUT6 and FUT7 genes are not expressed in erythroid cells.<sup>c</sup>Including hybrid alleles with another member of the GYPA/B/E gene family.

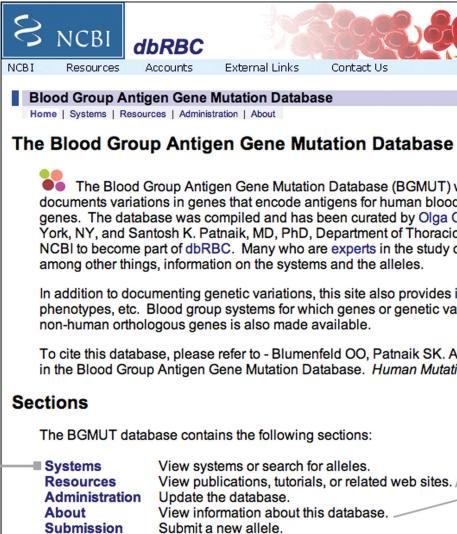
regions or ethnic populations and association of the allele with diseases are also noted.

Often, a name is also provided for an allele. Names make it easier to refer to alleles and can indicate the associated phenotype and/or nucleotide or amino acid variation. BGMUT uses the allele names given by the discoverers of the alleles and/or those that are commonly used by blood group system experts. In the absence of such a name, the curator may provide an arbitrary name. It should be noted that every allele entry in BGMUT has a unique database identification number associated with it.

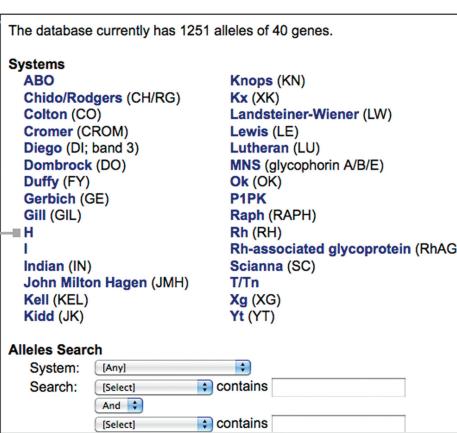
The naming of alleles has presented a problem compounded by names given to serological phenotypes of antigens they encode. At this time, efforts by the transfusion medicine community are in progress to standardize and simplify this nomenclature (6). In the database, to

more generally identify each allele, in most cases, ‘name’ includes the gene name followed by the DNA change particular to the allele; to facilitate the recognition of the phenotype/genotype connection, under ‘alias’ is given the serological designation as used by the original authors. In some cases, such as for the Rh system, because of traditional reasons, this arrangement is reversed; in others, such as ABO, only the phenotype designation marked by consecutive numbers is given; this is because the large number of sites of DNA changes in many alleles makes their listing under ‘name’ unwieldy.

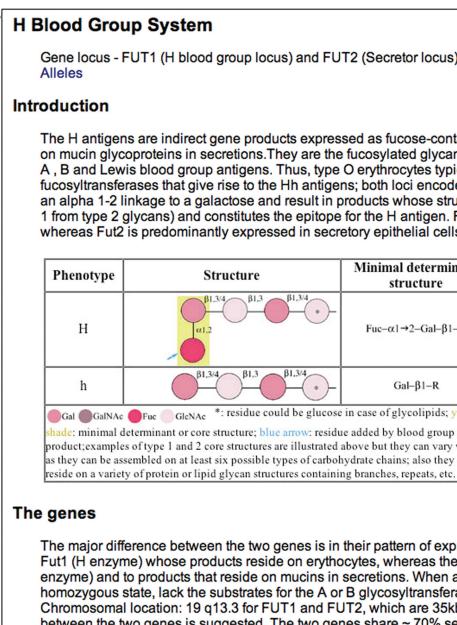
Alignments of DNA changes in alleles of a single gene or a gene family may uncover patterns that provide some insights into the process of diversification, including gene rearrangements and meiotic or unequal recombination (7). An attempt to correlate DNA changes in the gene(s) encoding the antigen(s), with the serological phenotype



The BGMUT home page features the NCBI dbRBC logo and a search bar. The main content area is titled "The Blood Group Antigen Gene Mutation Database" and includes sections for "publications", "tutorials", "related web sites", "database summary", "contributors", "help", "terminology", and "contact information".



The "Main systems page" lists 40 blood group systems: ABO, Chido/Rodgers (CH/RG), Colton (CO), Cromer (CROM), Diego (Di; band 3), Dombrock (DO), Duffy (FY), Gerbich (GE), Gill (GIL), H, I, Indian (IN), John Milton Hagen (JMH), Kell (KEL), Kidd (JK), Knops (KN), Kx (XK), Landsteiner-Wiener (LW), Lewis (LE), Lutheran (LU), MNS (glycophorin A/B/E), Ok (OK), P1PK, Raph (RAPH), Rh (RH), Rh-associated glycoprotein (RhAG), Sciana (SC), T/Tn, Xg (XG), and Yt (YT). It also includes an "Alleles Search" form.



The "Page for the H blood group system" includes an "Introduction" section and a table comparing the H and h blood group systems. The table shows the phenotype, structure, and minimal determining structure for each. A legend defines symbols for Gal, GalNAc, Fuc, GlcNAc, and Glc. A note states that the \* residue could be glucose in case of glycolipids.

+ non-human orthologs  
+ other system-related information

+ proteins and functions  
+ gene expression  
+ disease association  
+ nature of allelic variations  
+ links to external databases  
+ list of contributors

**Figure 1.** A montage of screenshots of the web-based interface of BGMUT depicting the hierarchically ordered home page, the page listing the blood group systems in the database and the page for a specific blood group system (H). The nature of content of the pages is visible in the screenshots or is noted in the figure.

**Alleles of the H Blood Group System**

**Search Parameters**

System: H  
Search: [Select] contains \_\_\_\_\_  
And [Select] contains \_\_\_\_\_  
Sort by: Name Ascending then [optional] Ascending  
**Search**

Next Page >> Page 1 Go Download Data

Name / Alias	Gene	Nucleotide Change	Amino Acid Change	Phenotype	Authors Pub Med IDs	Genbank Accession Nrs.	Details
1 FUT1	FUT1	reference	reference		Larsen et al. 2118655	M35531	Details...
2 FUT1-35T H*01.01.03	FUT1	-35C>T	5'UTR; none	O; a-b-	Matzhold et al. 19572973	Not submitted	Details...
3 FUT1 1047C	FUT1	1047G>C	W349C	Bombay	Wagner and Fiegel 9122901	Not submitted	Details...
4 FUT1 235C	FUT1	235G>C	G79R	para-Bombay; H+weak	Zhu FM and Xu XG	FJ665616	Details...
5 FUT1 293T	FUT1	293C>T	T98M	H+weak; para-Bombay	Yan et al. 15847661	AY611628	Details...
6 FUT1 328A	FUT1	328G>A	A110T	H+weak; para-Bombay	Guo et al. 15476160	DQ321370	Details...
7 FUT1 349T Reunion Island	FUT1	349C>T	H117Y	H+weak; Reunion	Fernandez-Mateos et al. 9745152	Not submitted	Details...
8 FUT1 35T H*11.01.01	FUT1	35C>T	A12V	H+	Xu XG et al.; Matzhold et al. 16331565, 19572973	Not submitted	Details...
9 FUT1 35T, 269T	FUT1	35C>T; 269G>T	A12V; G90V	para-Bombay	Story et al. 17176328	Not submitted	Details...
10 FUT1 35T, 423X h8(China)	FUT1	35C>T, 423G>A	A12V; W141X	para-Bombay	Cai X. and Xiang D. not published as yet	HQ699894, HM584610	Details...
11 FUT1 422A	FUT1	422G>A	W141X	Bombay; H-	Story et al. 17176328	Not submitted	Details...
12 FUT1 424T h9 (China)	FUT1	424C>T	R142W	para-Bombay	Chi Q, Zhang A, Guo Y, J, and Ren B, C. not published as yet	HQ891007	Details...
13 FUT1 442T h4 (Japan)	FUT1	442G>T	D148Y	para-Bombay; H+weak	Kaneko et al. 9226185	AB004862	Details...
14 FUT1 460C	FUT1	460T>C	Y154H	para-Bombay;	YU et al.	Not	Details...

**Details of FUT1 293T**

Database ID: 765  
System: H  
Gene: FUT1  
Name: FUT1 293T  
Alias:  
Nucleotide change: 293C>T  
Amino acid change: T98M  
Regions examined:  
Phenotype: H+weak; para-Bombay  
Prevalence: rare  
PubMed IDs: 15847661  
GenBank Accession Nrs.: AY611628  
Authors: Yan et al.  
Citations: Transfusion 2005 45 725-730  
Comment:  
Submitted by: O O. Blumenfeld, curator  
Last Update: 2010-03-23 20:11:51.887

## Page listing alleles of the H blood group system

## Page for details of an allele of the H blood group system

Figure 2. Screenshots of the BGMUT web pages showing a partial list of alleles and a specific allele of the H blood group system.

or the glycosyltransferase activity, shows that even though variation in common alleles occurs within the same sequence stretches that define the common epitopic regions (4), in rare alleles, changes that affect the serological response are seen throughout the exons. The resulting amino acid alterations may cause altered folding that interferes with epitope presentation or gives rise to new epitopes. As expected, null phenotypes usually result from absence of an intact protein and are caused by nonsense mutations, insertions or deletions, or absence of the common epitope due to recombination. More generally, in the systems documented in the database, the phenotype reflects a particular sequence alteration be it the result of a single or multiple molecular events (e.g. the Rh neg, ABO O and MNS Sta alleles). Clearly, in contrast to many

global studies, data presented in BGMUT allow to focus on DNA changes in a single gene and a closer look at its resulting alleles, to ask how they came about and to consider the properties of the resulting proteins. This knowledge will only expand as complete sequence content of variant alleles become more readily available.

### OTHER RESOURCES WITHIN BGMUT

Besides the alleles of 30 human blood group systems, BGMUT also provides information on the systems themselves (Figure 1). Such information includes that on a system's genes, such as their sequences, chromosomal locations and the functions of proteins encoded by them. Links to database records elsewhere, such as the Online

Mendelian Inheritance in Man (OMIM) (8) and PubMed resources of NCBI, are provided. Brief descriptions of diseases associated with a gene's variations, if any, and a summary of the nature of the variations are also provided. Content for such pages for the various blood group systems has been contributed by various experts and is continually updated. The BGMUT website also has a few introductory tutorials on blood group systems and provides links to online resources on blood groups as well as on human gene mutations and polymorphisms in general. Information on the nature of the genes of the H, MNS, Rh and RhAG systems in non-humans, primarily primates and their allelic variations is also provided. BGMUT also makes available brief descriptions of other carbohydrate antigens such as the Tk and Sid (Sda) antigens on erythrocytes.

## ADDITIONAL TOOLS OF INTEREST AT THE PARENT dbRBC RESOURCE

Besides BGMUT, the dbRBC resource contains tools and other resources developed at the NCBI to provide access to publicly available information on human blood group systems. Two of the tools are briefly described here. The alignment viewer tool (<http://www.ncbi.nlm.nih.gov/projects/gv/rbc/align.fcgi?cmd=aligndisplay>) provides an interface to choose alleles of genes of many of the blood group systems for sequence alignment, though allelic entries in BGMUT cannot be directly chosen in the current interface. Chosen sequences can be displayed as genomic DNA, cDNA or amino acid sequences. Positions known to bear gene polymorphisms can be highlighted, and so can be specific differences within a set of allelic sequences. The reference sequence itself can be chosen. Besides the alignment, a user can switch to a display of selected sequences in FASTA format. Nucleotide or protein sequences for individual exons and/or introns can be displayed for each allele in alignment to the reference sequence. When the three dimensional structure of a gene's protein product is known, amino acid differences of various alleles can be projected onto the protein model for viewing with the CN3D model viewer (9).

dbRBC's sequencing-based typing tool (<http://www.ncbi.nlm.nih.gov/projects/gv/rbc/sbt.cgi?cmd=main>) is of use to evaluate the allelic composition of sequencing-based typing results of cDNA or genomic sequences (10). It allows for the comparison of result sequences with sequences of known alleles of genes of various human blood group systems. The results of the comparison are returned as a table of potential allele hits, along with the respective nucleotide changes, and links for closer examination of the alignments within the dbRBC alignment viewer tool.

With a continuous increase in the number of available alleles, dbRBC is testing the implementation of an allelic nomenclature system based on one in use for the HLA system (11). Thus, for example, the reference *ABO* gene allele sequence is named *ABO*\*A1.01.01.01. The name consists of the gene name, followed by an asterisk, and for blocks separated by periods. The leading block is to group sequences first by their main serological phenotypes

(e.g. A1, A2 or Ax phenotypes of the ABO system). Sequences that share the same serological phenotype, but differ in their amino acid sequences, are differentiated by the second block. The third block lists alleles that have an identical amino acid sequence, but show synonymous mutations in their exonic DNA sequence. The last block differentiates sequences that share an identical DNA sequence in the coding region, but differ in non-coding regions. Alleles of the *ABO*, *FUT1*, *FUT2* and *FUT3* genes have been given additional names as per this system as of July 2011.

## FUTURE PLANS

Currently, allelic information compiled in BGMUT is not integrated with that outside the BGMUT but within the dbRBC resource. The latter set of alleles of genes for human blood group systems have been programmatically collected from various NCBI resources. Plans for an integration so that BGMUT allelic entries are available for use within the alignment viewer and the sequencing-based typing tools of dbRBC are under consideration. As a part of this integration, BGMUT alleles will be assigned additional names as per the nomenclature system outlined in the above section if a decision is made to implement it after consultation with various experts.

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## REFERENCES

- Sayers,E.W., Barrett,T., Benson,D.A., Bolton,E., Bryant,S.H., Canese,K., Chetvernin,V., Church,D.M., DiCuccio,M., Federhen,S. *et al.* (2011) Database resources of the National Center for Biotechnology Information. *Nucleic Acids Res.*, **39**, D38–D51.
- Broadberry,R.E. and Lin,M. (1996) The distribution of the MiiII (Gp.Mur) phenotype among the population of Taiwan. *Transfus. Med.*, **6**, 145–148.
- Claustres,M., Horaitis,O., Vanevski,M. and Cotton,R.G. (2002) Time for a unified system of mutation description and reporting: a review of locus-specific mutation databases. *Genome Res.*, **12**, 680–688.
- Blumenfeld,O.O. and Patnaik,S.K. (2004) Allelic genes of blood group antigens: a source of human mutations and eSNPs documented in the Blood Group Antigen Gene Mutation Database. *Hum. Mutat.*, **23**, 8–16.

5. Giardine,B., Riemer,C., Hefferon,T., Thomas,D., Hsu,F., Zielenki,J., Sang,Y., Elnitski,L., Cutting,G., Trumbower,H. *et al.* (2007) PhenCode: connecting ENCODE data with mutations and phenotype. *Hum. Mutat.*, **28**, 554–562.
6. Storry,J.R., Castilho,L., Daniels,G., Flegel,W.A., Garratty,G., Francis,C.L., Moulds,J.M., Moulds,J.J., Olsson,M.L., Poole,J. *et al.* (2011) International Society of Blood Transfusion Working Party on red cell immunogenetics and blood group terminology: Berlin report. *Vox Sang.*, **101**, 77–82.
7. Patnaik,S.K. and Blumenfeld,O.O. (2011) Patterns of human genetic variation inferred from comparative analysis of allelic mutations in blood group antigen genes. *Hum. Mutat.*, **32**, 263–271.
8. Hamosh,A., Scott,A.F., Amberger,J.S., Bocchini,C.A. and McKusick,V.A. (2005) Online Mendelian Inheritance in Man (OMIM), a knowledgebase of human genes and genetic disorders. *Nucleic Acids Res.*, **33**, D514–D517.
9. Wang,Y., Geer,L.Y., Chappay,C., Kans,J.A. and Bryant,S.H. (2000) Cn3D: sequence and structure views for Entrez. *Trends Biochem. Sci.*, **25**, 300–302.
10. Helmberg,W., Dunivin,R. and Feolo,M. (2004) The sequencing-based typing tool of dbMHC: typing highly polymorphic gene sequences. *Nucleic Acids Res.*, **32**, W173–W175.
11. Robinson,J., Mistry,K., McWilliam,H., Lopez,R., Parham,P. and Marsh,S.G. (2011) The IMGT/HLA database. *Nucleic Acids Res.*, **39**, D1171–D1176.